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(A) HIGH PERFORMANCE PRESSURE-SENSITIVE ADHESIVE POLYMERS.

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BACKGROUND OF THE INVENTION

The present invention is directed to pressure-sensitive adhesive compositions that exhibit high adhesion to high energy surfaces such as aluminum and possess high tack and outstanding holding power at elevated temperatures. Two key monomers are used in combination in the pressure-sensitive adhesives of this invention to impart unique properties to the resulting polymers. The monomers are at least one glycidyl monomer in combination with at least one N-vinyl lactam.

The use of glycidyl monomers in pressure-sensitive adhesive has been disclosed in the art.

U.S. Patent 3,284,423 discloses creep-resistant pressure-sensitive adhesive compositions comprising 35-75% by weight alkyl acrylate esters containing 6-15 carbon atoms, 10-60% lower alkyl acrylate, 0.1-10% by weight of an ethylenically unsaturated carboxylic acid and 0.1-10% by weight glycidyl ester.

U.S. Patent 3,893,982 discloses an interpolymer comprising 0.1-15% parts of an ethylenically unsaturated carboxylic acid, 0.1-2% parts of a glycidyl monomer, 35-84.9% parts of an alkyl acrylate or methacrylate and optionally a monomer selected from the group consisting of alpha-olefins containing 2-10 carbon atoms, vinyl esters of alkanoic acids containing 3-10 carbon atoms, ethyl and methyl esters of acrylic and methacrylic acids, acrylonitrile, methacrylonitrile, styrene and vinyl chloride where the polymer has a weight average molecular weight in the range of 10,000 - 500,000 and between 0.01 and 1 parts by weight per 100 parts of the copolymer of 1,3-bis(dimethylamino)-2-hydroxypropane to cause the cure of the epoxy group.

The art has also taught the use of N-vinyl lactams in polymers.

U.S. Patent 3,728,148 discloses a pressure-sensitive adhesive for electrical insulating applications comprising of a copolymer of 65-90% by weight of an alkyl acrylate ester, 10-30 by weight of a N-vinyl lactam and 0-20% by weight of a modifying monomer which is copolymerizable with the above. Acidic monomers and amides are excluded as they are claimed to cause undesirable corrosion.

U.S. Patent 4,181,752 discloses an interpolymer containing 87% by weight isooctyl acrylate, 8% by weight vinyl pyrrolidone, 3% by weight acrylic acid and 2% by weight acrylamide.

U.S. Patent 4,364,972 discloses a pressure-sensitive adhesive tape made by copolymerizing an alkyl acrylate ester with 15 to 50 parts by weight of vinyl pyrrolidone and having a K-value greater than 100 and when crosslinked has a gel-swell in ethyl acetate in excess of 600%. Advantages claimed are good adhesion to automotive paints, rubber and plastic foam layers.

U.S. Patent 4,310,509 discloses a 90/10 2-ethyl hexyl acrylate/vinyl pyrrolidone copolymer for making a pressure sensitive adherent for complexing with iodine for anti-microbial activity.

European Patent Application 130080 discloses an emulsion polymerized pressure-sensitive adhesive comprising 2-20% by weight N-vinyl lactam and an alkyl acrylate ester. The claimed use is good adhesion to skin under hot and humid conditions.

U.S. Patent 4,370,380 is directed to a blend of two polymers. One is a copolymer of 88-99% by weight of an alkyl acrylate ester with 1-12% by weight of a carboxylic acid with a glass transition temperature (Tg) of less than 0°C. The second polymer is either a homo or a copolymer of N-vinyl lactam with a Tg of 20-150°C. The blend ratio is 70-99% by weight of the first polymer with 1-30% by weight of the second polymer. The resultant pressure-sensitive adhesive is disclosed to have moisture permeability.

U.S. Patent 4,150,197 discloses a water vapor permeable pressure-sensitive adhesive comprising a copolymer of 79-89% of butyl acrylate, 10-20% by weight of N-vinyl lactam and 1-5% by weight of an acidic comonomer.

None of the patents or applications discloses a copolymer containing both a glycidyl monomer and N-vinyl lactam monomer.

SUMMARY OF THE INVENTION

It has now been found that, as part of an acrylic and/or methacrylic ester based polymer system, a glycidyl monomer and a N-vinyl lactam monomer synergistically act to provide unusually high adhesion to high energy surfaces such as aluminum and stainless st el. The pr ssure-sensitiv adhesiv s of the invintion are formed of copolymers containing essentially no cross-linking when polymeriz d and which contain on a copolym rized basis from about 0.01 to about 2% by w ight of glycidyl monomer, about 1% t about 20% by weight, pr f rably from about 1 to about 10% by weight of an N-vinyl lactam monom r, from 0 to about 15% by w ight of an thylenically unsaturated carboxylic acid, from about 55 to about 85% by w ight an alkyl acrylate or methacrylate ester containing fr m 4 to 12 carbon atoms in th alkyl group, from 0 to 35% by weight f an alkyl acrylate or methacrylat ester containing less than 4 carbon atoms in the alkyl group, and optionally, one or more other mon mers mpl yed to tailor polymer properties, such as glass transition temperatur, to end use applications. Such monomers includ polystyryl thyl methacrylate, acetoacetoxy ethyl methacrylate, styrene, alpha olefins, and vinyl est rs of alkanoic acids containing greater than 3 carbon atoms and mixtures thereof. Modifying

monomer content can range from 0 to ab ut 35% by weight of the total monomers.

The ratio of monomers is selected to provide a copolymer with glass transition temperature of less than about -15°C and a weight average molecular weight of at least about 200,000, preferably from about 200,000 to about 500,000 as determined by size exclusion chromatography using polystyrene for calibration. Polymers of the instant invention may be synthesized by solution, emulsion and bulk polymerization. It is presently preferred that they be formed by solution polymerization. Polymers are cross-linked to the desired extent, prior to use, using heat, an ionic additives, actinic or electron beam radiation.

The polymers on cross-linking exhibit excellent adhesion to high energy surfaces such as aluminum as evidenced by increased peel adhesion with dwell and superior rivet performance as evidenced by reduced tenting as described herein.

THE DRAWINGS: 15

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FIG. 1 graphically compares 180° peel adhesion on alodine aluminum panels of two adhesives, one which contains both a glycidyl monomer and a lactam monomer to one containing only a glycidyl monomer.

FIG. 2 compares 180° peel on stainless steel as a function of dwell at two temperature conditions, room temperature and 120°C for compositions of the invention to controls.

The room temperature dwell was for 20 minutes and 120°C dwell was for 30 minutes. After dwelling for 30 minutes at 120°C, the adhesive on the substrate was allowed to equilibrate to room temperature and 180° peel determined.

DETAILED DESCRIPTION

Pressure-sensitive adhesive polymers of the instant invention are prepared by copolymerizing a mixture of monomers comprising from about 55 to about 85% by weight of an alkyl acrylate and/or methacrylate ester containing 4 to about 12 carbon atoms in the alkyl group; from about 0.01 to about 2% by weight of a glycidyl monomer, from about 1 to about 20% by weight, preferably from 1 to about 10% by weight of an N-vinyl lactam monomer; from 0 to 15% by weight, preferably from about 5 to about 13% by weight of an unsaturated carboxylic acid; from 0 to about 35% by weight of an alkyl acrylate and/or methacrylate ester containing less than 4 carbon atoms in the alkyl group and optionally from about 0 to 35% by weight of one or more other comonomers to provide a balance of desirable polymer properties such as glass transition temperature. The precise ratio of the monomers is selected to give a polymer whose glass transition temperature is lower than about -15°C. The polymers of the instant invention have a weight average molecular weight of at least about 200,000, preferably from about 200,000 to about 500,000 as determined by size exclusion chromatography using polystyrene as the calibrator.

The alkyl acrylate and methacrylate esters containing 4 to 12 carbon atoms in the alkyl group useful in forming the polymers of the instant invention include without limitation 2-ethyl hexyl acrylate, isooctyl acrylate, butyl acrylate, sec-butyl acrylate, methyl butyl acrylate, 4-methyl-2-pentyl acrylate, isodecyl methacrylate and mixtures thereof. Isooctyl acrylate and 2-ethyl hexyl acrylate are presently preferred.

The glycidyl monomers are glycidyl acrylate, glycidyl methacrylate, allyl glycidyl ether and mixtures thereof. The presently preferred gylcidyl monomer is glycidyl methacrylate.

The N-vinyl lactams monomers which may be used include N-vinyl pyrrolidone, N-vinyl caprolactam, 1-vinyl-2-piperidone, 1-vinyl-5-methyl-2-pyrrolidone, N-vinyl pyrrolidone is presently preferred.

Ethylenically unsaturated carboxylic acids include acrylic acid, methacrylic acid, fumaric acid.

Alkyl acrylate and methacrylate esters containing less than 4 carbon atoms in the alkyl group include methyl acrylate, ethyl acrylate, methyl methacrylate. Methyl acrylate is presently preferred.

Other monomers which can be included are polystyryl ethyl methacrylate, acetoacetoxy ethyl methacrylate, alpha olefins such as ethylene and propylene and vinyl esters of alkanoic acids containing more than three carbon atoms as well as mixtures th reof. Such monomer concentrations are in the range from 0 to about 35 percent by w ight of the total monom rs. A preferred pressure-sensitive adhesive is described in claim 7.

The copolym rs may be synthesiz d using solution mulsion, and batch polymerization t chniques. It is presently pref rr d to prepare the copolymers in solution using a mixture of solv nts. Th present preferr d solution polym rization involv s the us of bl nds of thyl acetat and hexane or thyl acetate and acetone. The ratio of solv nts are adjusted to provide a reflux temp rature of from about 68°C to ab ut 78°C. Solids cont nt during polymerization may typically range from about 40% to about 60% in order to achieve the desired w ight average molecular w ight, and yet achieve viscosities that ar manageable in the reactor. R action occurs in the presence of fire-radical initiators, preferably of the azotype, for example, 2,2'-aze bis (isobutyronitrile). The polymers form of are solvent soluble polymers with essentially no crosslinking. To this ond, the glycidyl monomer is preferably limited to 2% by weight of the total monomers to avoid the possibility of crosslinking, by opening of the oxirane group, during polymerization or during aging. Polymers can, as desired, be post-polymerization cross-linked using heat, actinic or electron beam radiation.

The unique characteristics of the cross-linked pressure-sensitive adhesive copolymers of the instant invention is a dramatic adhesion to high energy surfaces, such to aluminum and stainless steel, as seen by increased peel adhesion values with dwell times and superior rivet performance, as reflected by reduced tenting. Superior performance requires the presence of both glycidyl and lactam monomers. In the absence of one, poor adhesion to aluminum has been observed, as manifested by poor tenting performance (see Table I). These results are unexpected as a primary purpose in incorporating the N-vinyl lactam is to increase adhesion to vinyl films and painted surfaces. The purpose of the glycidyl group is normally to introduce a latent functional group which could undergo cross-linking at elevated temperatures under use conditions. This was realized by the dramatic improvement in elevated temperature performance even as high as 200°C and high levels of shear adhesion failure temperature (SAFT) of polymers containing glycidyl methacrylate as established by Table II. Improved peel adhesion was totally unexpected. While not bound by theory, it is presently believed that chemical reaction occurs between the epoxy groups and functional groups such as hydroxyl groups on the substrate or a complexation reaction occurs between the epoxy groups and the nitrogen of the lactam, with ionic groups such as A1+++ on the substrate leading to increased bond strength.

An application of the adhesives of the instant invention is the marking of truck panels and the like with an adhesive coated vinyl film. The film is applied over a panel which may be a painted or unpainted aluminum panel fixed with aluminum rivets. The ability of the adhesive to conform to the contour of the rivet and not lift appreciably after application is highly desirable. A tendency to lift away is known as "tenting" and the greater the separation from the rivet the more unsatisfactory is the adhesive.

The SAFT test is a test where the adhesive is applied to 0.5" x 1" overlap on stainless steel to which a 4.5 lb. roll force applied. After dwell of 24 hours, this is placed in an oven and a kilogram load is applied under shear conditions and temperature raised from 40°C to 200°C at the rate of 1°C per minute. The failure temperature is recorded as the shear adhesion failure temperature. This is a measure of the cohesive strength of the adhesive or the ability of the adhesive to maintain a bond at elevated temperatures.

While not limiting, the following illustrate the invention.

Example 1

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A monomer mixture was made up by mixing 423 g. of 2-ethyl hexyl acrylate, 145 g of methyl acrylate, 3.15 g. of glycidyl methacrylate, 12.6 g of N-vinyl pyrrolidone and 44.1 g of acrylic acid. 157 g. of this mixture was introduced to a 2 liter reactor equipped with a pitched turbine agitator, a reflux condensor and a thermistor. Also added were 73.5 g. of ethyl acetate and 78.76 g of hexane. The contents of the reactor were heated to reflux and 0.238 g of Vazo 64, manufactured and sold by duPont in 5.0 g of ethyl acetate was added. Vigorous reflux started in a short time and the contents of the reactor were held for 23 minutes. At this time, the remaining monomers were mixed with 537.2 g. of ethyl acetate, 75.2 g. of hexane and 0.707 g. of Vazo 64 and added as a single feed mixture over 3.5 hrs. All through the feed, temperature was maintained to keep reactor contents under reflux. One hour after end of feed, 0.17 g. Vazo 64 was added in 5 g. ethyl acetate and held for an additional hour. The percentage of solids content at the end of reaction was 46.4% and the viscosity was 23 Pa.s using #4 @ 12 on a Brookfield viscometer.

Example 2

A monomer mixture was made up by mixing 453.6 g. of Isooctyl acrylate, 100.8 g of methyl acrylate, 6.3 g. of glycidyl methacrylate, 25.2 g of N-vinyl caprolactam, 44.1 g of acrylic acid and 0.945 g of Vazo 64. 157.5 g of this mixture was introduced to the reactor with 78.76 g of hexane, 78.76 g of ethyl acetate and heated to reflux. Once vigorous reflux initiat d, the contents were held for about 12 minutes and the remaining monomers add d along with 537.24 g of ethyl acetate and 75.24 g of hexan as a single feed over 3 hours. Two hours after the nd of feed, the contents were cooled.

Example 3 - Tenting T st

There was formed as control 1 a polymer containing 65.7% by w ight 2-ethyl hexyl acrylate, 27.3% by

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w ight m thyl acrylate and 7% by w ight acrylic acid. As Control 2, th re was formed a polymer containing 67% by weight 2-ethyl h xyl acrylate, 24% by weight methyl acrylate, 7% acrylic acid, and 2% by w ight N-vinyl pyrrolidone. As Control 3, there was form dap lymer containing 65.7% by weight 2-ethyl hexyl acrylate, 27.2% by weight methyl acrylate, 7% by weight acrylic acid and 0.1% by weight glycidyl methacrylate. As Control 4, there was used a polymer containing 65% by weight 2-ethyl hexyl acrylate, 27% by weight methyl acrylate, 7% by weight acrylic acid and 1% by weight glycidyl methacrylate. These were compared for tenting in a rivet test to the polymers of Examples 1 and 2. As a cross-linker in each instance, there was added 0.2 parts by weight of the polymer of aluminum acetyl acetonate. For the rivet tenting test adhesive was transfer coated from a release liner to a cast vinyl facestock, at a coat weight of 30 g/m². The results are given in Table 1, wherein the lower the value reported, the less tenting, i.e., lifting away from the rivet, occurred.

Table 2 compares the polymer of Example 1 electron-beam (EB) cured at a dosage of 30 kiloGray (kGy) and the polymer of Example 2 also EB cured at a dosage of 30 kGy. Examples 1 and 2 exhibited the best combinations of shear and tack.

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TABLE 1

5	Control/ Example*	Tenting in Rivet Test, mm
	Control 1	1.42
10	Control 2	1.32
	Control 3	1.37
	Control 4	1.25
15	Example 1	0.58
	Example 2	1.1

* All the polymers were cross-linked with 0.2 parts per hundred parts of the polymer of Aluminum acetyl acetonate.

TABLE 2

25		Example 1	Example 2
	180° Peel, NM	EB, 30 kGy	EB, 30 kGy
30	Stainless Steel, 20' Dwell	572	632
	Hard PVC	692	780
	Static Shear, min.		
35	20°C/1kg	10000+	9577
	150°C/1kg	5200+	5200+
40	200°C/1kg	3300+	3300+
40	Loop Tack, N/M	600	270
	SAFT, 'C	200*+	200*+
			

RT Shear, 0.5 x 0.5 inch overlap, Al Facestock SAFT and ET Shear, 0.5 x 1.0 inch overlap, Al Facestock For SAFT, 24 hr. dwell, rate of heating 1°C/minute

50 Control 5 to 8 and Examples 4 and 5

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There was compared to the product of Example 2 a copolymer containing 67 parts by weight isooctyl acrylate, 25 parts by weight methyl acrylate, 1 part by weight glycidyl methacrylate, and 7 parts by weight acrylic acid (Control 5). Control 5 and Exampl 2 copolymers were coated to a level of 50 g/m² ont mylar and cured at an 1 ctron b am lev 1 of 30 kGy. A comparison of 180° peel on alodine aluminum substrat in Newtons-/meter (N/M) as a function of dw II time is shown in Figure 1 and establishes that the combination of glycidyl methacrylate and vinyl caprolactam produce unusually high 180° peels, especially after extended dwell times.

Figure 2 compar s 180° peel on stainless steel substrate as a function of dwell at room temperature for

20 minutes and dwell at 120°C for 30 minut s followed by peel testing at room t mperatur. All polym rs employed contained 2-ethyl h xyl acrylat , m thyl acrylate and acrylic acid. Control 6 contained no glycidyl methacrylate or N-vinyl lactam. Control 7 contained 0.1 part glycidyl m thacrylate but no N-vinyl lactam. Control 8 contained 1 part by weight glycidyl methacrylate but no N-vinyl lactam. Example 4 contained 0.1 part by weight glycidyl methacrylate and 2 parts by weight N-vinyl pyrrolidone. Example 5 contained 0.1 part by weight glycidyl methacrylate and 4 parts by weight N-vinyl caprolactam per hundred parts total monomer. In each instance, the polymers were EB cured at a dosage of 30 kGy. The glycidyl methacrylate in combination and N-vinyl lactam gives better performance in terms of adhesion to high energy surfaces than the individual constituents of the combination.

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Claims

- 1. A pressure-sensitive adhesive comprising a copolymer comprising on a copolymerized basis from about 55 to about 85% by weight of a monomer selected from the group consisting of alkyl acrylate esters and alkyl methacrylate esters containing from 4 to 12 carbon atoms in the alkyl group and mixtures thereof, from 0 to about 35% by weight of an alkyl acrylate or methacrylate ester containing less than 4 carbon atoms in the alkyl group, from 0.01 to about 2% by weight of a glycidyl monomer, from about 1 to about 10% by weight of an N-vinyl lactam, and from 0 to about 15% by weight of an unsaturated carboxylic acid, said copolymer having a weight average molecular weight of at least about 200,000 and a glass transition temperature less than about -15°C.
- 2. A pressure-sensitive adhesive as claimed in claim 1 which contains, based on the total weight of monomers, up to about 35% by weight, of a monomer selected from the group consisting of polystyryl ethyl methacrylate, aceto-acetoxy ethyl methacrylate, styrene, alpha olefins, vinyl esters of alkanoic acids containing more than three carbon atoms and mixtures thereof.
- 3. A pressure-sensitive adhesive as claimed in claim 1 or 2 in which the alkyl acrylate is 2-ethyl hexyl acrylate or isooctyl acrylate.
- 4. A pressure-sensitive adhesive as claimed in any one of claims 1 to 3 in which the glycidyl monomer is selected from the group consisting of glycidyl acrylate, glycidyl methacrylate, allyl glycidyl ether and mixtures thereof.
- A pressure-sensitive adhesive as claimed in any one of claims 1 to 4 in which the N-vinyl lactam is selected from the group consisting of N-vinyl pyrrolidone, N-vinyl caprolactam and mixtures thereof.
- 6. A pressure-sensitive adhesive as claimed in any one of claims 1 to 5 in which the formed copolymer is cross-linked by exposure to heat, anionic additive, actinic radiation or electron beam radiation.
- 7. A pressure-sensitive adhesive comprising a copolymer comprising from about 55 to about 85% of an alkyl acrylate ester selected from the group consisting of isooctyl acrylate and 2-ethyl hexyl acrylate, from about 0.01 to about 2% by weight glycidyl methacrylate, from about 1 to about 10% of a N-vinyl lactam selected from the group consisting of N-vinyl pyrrolidone and N-vinyl caprolactam, from about 5 to 13% by weight unsaturated carboxylic acid, and up to about 35% by weight methyl acrylate, said polymer having weight average molecular weight from about 200,000 to about 500,000 and a glass transition temperature less than about -15°C.
- 8. A pressure-sensitive adhesive as claimed in claim 18 in which the formed copolymer is cross-linked by exposure to heat, anionic additive, actinic radiation or electron beam radiation.

Patentansprüche

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- 1. Haftkleber, bestehend aus einem Copolymeren, das auf Copolymerisationsbasis etwa 55 bis etwa 85 Gew.-% eines Monomeren, das aus der aus Alkylacrylatestern und Alkylmethacrylatestern mit 4 bis 12 Kohlenstoffatomen in der Alkylgruppe und Mischungen davon bestehenden Gruppe ausgewählt ist, 0 bis etwa 35 Gew.-% eines Alkylacrylat oder Methacrylatesters mit weniger als 4 Kohlenstoffatomen in der Alkylgruppe, 0,01 bis etwa 2 Gew.-% eines Glycidylmomeren, etwa 1 bis 10 Gew.-% eines N-Vinyllactams und 0 bis etwa 15 Gew.-% einer ungesättigten Carbonsäure enthält, wobei das Copolymere ein Gewichtsmittel-Molekulargewicht von zumindest etwa 200.000 und eine Glasübergangstemperatur von w niger als etwa -15°C besitzt.
- 2. Haftkleber nach Anspruch 1, der, auf der Basis des Gesamtg wichts der Monomeren, bis zu etwa 35 Gew.-% eines M nomer n nthält, das aus der aus Polystyryl thylmethacrylat, Aceto-Acetoxyethylmethacrylat, Styrol, Alphaolefin n, Vinylest rn von Alkansäur n mit mehr als drei Kohl nstoffatom n und Mischungen dav n best henden Gruppe ausg wählt ist.
- 3. Haftkleber nach Anspruch 1 oder 2, bei dem das Alkylacrylat ein 2-Ethylhexylacrylat od r Isooctylacrylat ist.

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- 4. Haftkleber nach einem der Ansprüche 1 bis 3, b i dem das Glycidylmonom re aus der aus Glycidylacrylat, Glycidylmethalcrylat, Allylglycidyl th r und Mischung n davon b steh nden Grupp ausg wählt ist.
- 5. Haftkleb r nach einem der Ansprüche 1 bis 4, bei dem das N-Vinyllactam aus der aus N-Vinylpyrrolidon, N-Vinylcaprolactam und Mischungen davon bestehenden Gruppe ausg wählt ist.
- 6. Haftkleber nach einem der Ansprüche 1 bis 5, bei dem das geformte Copolymere durch Aussetzen der Wärme, einem anionischen Zusatz, einer aktinischen Strahlung oder einer Strahlung mittels Elektronenstrahl vernetzt ist.
- 7. Haftkleber, bestehend aus einem Copolymeren, das etwa 55 bis etwa 85% eines Alkylacrylatesters, der aus der aus Isooctylacrylat und 2-Ethylhexylacrylat bestehenden Gruppe ausgewählt ist, etwa 0,01 bis etwa 2 Gew.-% Glycidylmethacrylat, etwa 1 bis etwa 10% N-Vinyllactam, das aus der aus N-Vinylpyrrolidon und N-Vinylcaprolactam bestehenden Gruppe ausgewählt ist, etwa 5 bis 13 Gew.-% ungesättigter Carbonsäure und bis zu etwa 35 Gew.-% Methylacrylat enthält, wobei das Polymere ein Gewichtsmittelmolekularyewicht von etwa 200.000 bis etwa 500.000 und eine Glasübergangstemperatur von weniger als etwa -15°C besitzt.
- 8. Haftkleber nach Anspruch 7, bei dem das geformte Polymere durch Aussetzen der Wärme, einem anionischen Zusatz, einer aktinischen Strahlung oder einer Strahlung mittels Elektronenstrahl vernetzt ist.

Revendications

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- 1. Adhésif sensible à la pression, comprenant un copolymère composé, à l'état copolymérisé, d'environ 55 à environ 85% en poids d'un monomère choisi dans le groupe constitué par les esters acrylates d'alkyles et les esters méthacrylates d'alkyles contenant de 4 à 12 atomes de carbone dans le groupement alkyle et par leurs mélanges, de 0 à environ 35% en poids d'un ester acrylate ou méthacrylate d'alkyle contenant moins de 4 atomes de carbone dans le groupement alkyle, de 0,01 à environ 2% en poids d'un monomère glycidylique, d'environ 1 à environ 10% en poids d'un N-vinyl-lactame et de 0 à environ 15% en poids d'un acide carboxylique insaturé, ledit copolymère ayant un poids moléculaire moyen pondéral d'au moins 200 000 environ et une température de transition vitreuse inférieure à -15°C environ.
- 2. Adhésif sensible à la pression selon la revendication 1, contenant, sur la base du poids total des monomères, jusqu'à 35% en poids environ d'un monomère choisi dans le groupe constitué par le méthacrylate de polystyryléthyle, le méthacrylate d'acétylacétoxyéthyle, le styrène, les α-oléfines, les esters vinyliques d'acides alcanoïques contenant plus de 3 atomes de carbone et les mélanges de ces monomères.
- 3. Adhésif sensible à la pression selon la revendication 1 ou 2, dans lequel l'acrylate d'alkyle est l'acrylate de 2-éthylhexyle ou l'acrylate d'isoocytyle.
- 4. Adhésif sensible à la pression selon l'une quelconque des revendications 1 à 3, dans lequel le monomère glycidylique est choisi dans le groupe constitué par l'acrylate de glycidyle, le méthacrylate de glycidyle, l'allylglycidyléther et les mélanges de ces monomères.
- 5. Adhésif sensible à la pression selon l'une quelconque des revendications 1 à 4, dans lequel le N-vinyllactame est choisi dans le groupe constitué par la N-vinylpyrrolidone, le N-vinylcaprolactame et les mélanges de ceux-ci.
- 6. Adhésif sensible à la pression selon l'une quelconque des revendications 1 à 5, dans lequel le copolymère formé est réticulé par exposition à la chaleur, à un additif anionique, à un rayonnement actinique ou à un faisceau électronique.
- 7. Adhésif sensible à la pression, comprenant un copolymère qui comprend environ 55 à environ 85% d'un ester acrylate d'alkyle choisi dans le groupe constitué par l'acrylate d'isooctyle et l'acrylate de 2-éthylhexyle, environ 0,01 à environ 2% en poids de méthacrylate de glycidyle, environ 1 à environ 10% d'un N-vinyl-lactame choisi dans le groupe constitué par la N-vinylpyrrolidone et le N-vinylcaprolactame, environ 5 à 13% en poids d'acide carboxylique insaturé et jusqu'à 35% en poids environ d'acrylate de méthyle, ledit polymère ayant un poids moléculaire moyen pondéral d'environ 200 000 à environ 500 000 et une température de transition vitreuse inférieure à -15°C environ.
- 8. Adhésif sensible à la pression selon la revendication 7, dans lequel le copolymère formé est réticulé par exposition à la chaleur, à un additif anionique, à un rayonnement actinique ou à un faisceau électronique.

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